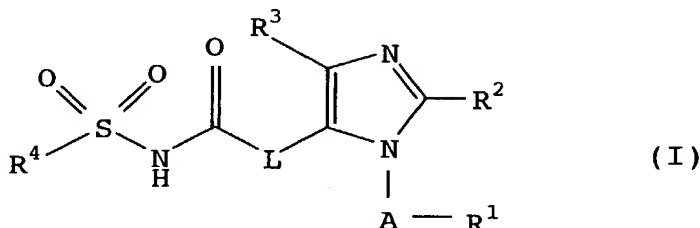


WHAT IS CLAIMED IS

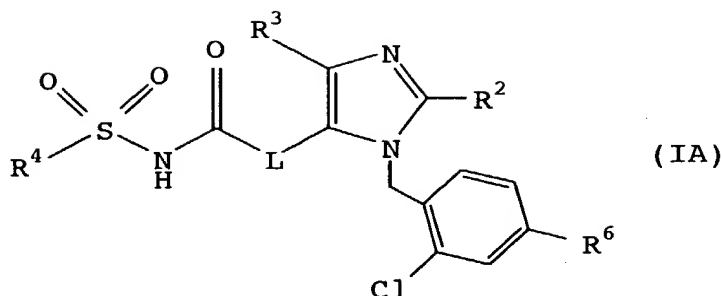
1. An imidazole compound of the formula (I):



5 wherein

- R^1 is an aryl or heterocyclic group substituted by a substituent selected from the group consisting of (1) aryl, (2) heterocyclic group, (3) halogen, (4) halo(lower)alkyl, (5) lower alkylthio, (6) nitro, (7) lower alkenyl optionally substituted by aryl, (8) lower alkynyl optionally substituted by aryl, (9) lower alkoxy optionally substituted by cyclo(lower)alkyl or aryl, (10) aryloxy and (11) amino optionally substituted by protected carboxy or lower alkyl;
- 15 R^2 is a lower alkyl;
- R^3 is a hydrogen, halogen, lower alkyl or nitro;
- R^4 is (1) a lower alkenyl optionally substituted by aryl or heterocyclic group, (2) aryl optionally substituted by lower alkenyl, (3) lower alkyl, or (4) heterocyclic group optionally substituted by halogen;
- 20 A is a lower alkylene; and
- L is a single bond, lower alkenylene or lower alkylene optionally substituted by aryl or heterocyclic group, or $-X-CH_2-$ wherein X is $-O-$, NR^5 wherein R^5 is hydrogen or lower alkyl, or $-S-$,
- 25 or a salt thereof.

2. The imidazole compound of claim 1, which has the formula (IA):



wherein

R^2 is methyl;

R^3 is chlorine;

R^4 is (1) lower alkenyl optionally substituted by aryl, (2) aryl, (3) lower alkyl, or (4) heterocyclic group optionally substituted by halogen;

R^6 is (1) aryl, (2) heterocyclic group, (3) bromine, (4) halo(lower)alkyl, (5) lower alkylthio, (6) nitro, (7) lower alkenyl substituted by aryl, (8) lower alkynyl substituted by aryl, (9) lower alkoxy optionally substituted by cyclo(lower)alkyl or aryl, (10) lower alkyl optionally substituted by aryloxy, or (11) amino optionally substituted by protected carboxy or lower alkyl; and

L is ethenylene,

or a salt thereof.

3. The imidazole compound of claim 2, wherein R^4 is aryl, or lower alkenyl optionally substituted by aryl, R^6 is bromine, lower alkenyl substituted by aryl, lower alkynyl substituted by aryl, or lower alkoxy optionally substituted by cyclo(lower)alkyl, or a salt thereof.

4. The imidazole compound of claim 1, wherein R^1 is heterocyclic group substituted by a substituent selected from the group consisting of (1) aryl, (2) heterocyclic group, (3) halogen, (4) halo(lower)alkyl, (5) lower alkylthio, (6) nitro, (7) lower alkenyl optionally substituted by aryl, (8) lower alkynyl optionally substituted by aryl, (9) lower alkoxy optionally substituted by cyclo(lower)alkyl or aryl, (10) aryloxy and (11) amino optionally substituted by protected carboxy or lower alkyl, or a salt thereof.

5. The imidazole compound of claim 1, which is:

- 40 methylimidazol-5-yl)-N-((E)-2-phenylethenyl)sulfonyl)-2-

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- (52) (E)-3-(4-chloro-1-(2-chloro-4-(1-pentyloxy)benzyl)-2-ethylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(53) (E)-3-(1-(4-bromo-2-chlorobenzyl)-4-chloro-2-ethylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
5 (54) (E)-3-(4-chloro-1-(2-chloro-4-(phenylethynyl)benzyl)-2-ethylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(55) (E)-3-(4-chloro-1-(2-chloro-4-(phenylethynyl)benzyl)-2-ethylimidazol-5-yl)-N-((4-methylbenzene)sulfonyl)-2-propenamide,
(56) (E)-N-(1-butanesulfonyl)-3-(4-chloro-1-(2-chloro-4-(phenylethynyl)benzyl)-2-ethylimidazol-5-yl)-2-propenamide,
10 (57) (E)-3-(4-chloro-1-(2-chloro-4-(phenylethynyl)benzyl)-2-ethylimidazol-5-yl)-N-(1-pentanesulfonyl)-2-propenamide,
(58) (E)-3-(4-chloro-1-(2-chloro-4-(phenylethynyl)benzyl)-2-ethylimidazol-5-yl)-N-((E)-1-penten-1-ylsulfonyl)-2-propenamide,
15 (59) (E)-3-(4-chloro-1-(2-chloro-4-((E)-2-phenylethenyl)benzyl)-2-ethylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(60) (E)-3-(4-chloro-1-(2-chloro-4-((E)-2-phenylethenyl)benzyl)-2-ethylimidazol-5-yl)-N-(4-methylbenzenesulfonyl)-2-propenamide,
(61) (E)-N-(1-butanesulfonyl)-3-(4-chloro-1-(2-chloro-4-((E)-2-phenylethenyl)benzyl)-2-ethylimidazol-5-yl)-2-propenamide,
20 (62) (E)-3-(4-chloro-1-(2-chloro-4-((E)-2-phenylethenyl)benzyl)-2-ethylimidazol-5-yl)-N-(1-pentanesulfonyl)-2-propenamide,
(63) (E)-3-(4-chloro-1-(2-chloro-4-((E)-2-phenylethenyl)benzyl)-2-ethylimidazol-5-yl)-N-((E)-1-penten-1-ylsulfonyl)-2-propenamide,
25 (64) (E)-3-(1-(4-bromo-2-chlorobenzyl)-2,4-dimethylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(65) (E)-3-(4-bromo-1-(2-chloro-4-(1-pentyloxy)benzyl)-2-methylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(66) (E)-3-(1-(2-chloro-4-(1-pentyloxy)benzyl)-4-ethyl-2-methylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
30 (67) (E)-2-benzyl-3-(1-(2-chloro-4-(1-pentyloxy)benzyl)-2-methylimidazol-5-yl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(68) (E)-3-(1-(2-chloro-4-(1-pentyloxy)benzyl)-2-methylimidazol-5-yl)-2-(1-pentyl)-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
35 (69) (E)-3-(1-(2-chloro-4-(1-pentyloxy)benzyl)-2-methylimidazol-5-yl)-2-(3-pyridyl)methyl-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
(70) (E)-3-(1-(2-chloro-4-(1-pentyloxy)benzyl)-2-methylimidazol-5-yl)-2-methyl-N-((E)-2-phenylethanesulfonyl)-2-propenamide,
40 (71) (E)-3-(4-chloro-1-(2-chloro-4-(1-pentyloxy)benzyl)-2-

hypertension, inflammatory bowel diseases, skin disorders related to an anomaly of differentiation of epidermic cells, angina pectoris, pulmonary hypertension, congestive heart failure, glomerulopathy, tubulointerstitial disorders, renal failure, angiostenosis, peripheral vascular diseases, cerebral apoplexy, chronic reversible obstructive impairment, autoimmune diseases, allergic rhinitis, urticaria, glaucoma, diseases characterized by impaired intestinal motility, impotence, nephritis, cancer cachexia, restenosis after PTCA, or cachexia.

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9. Use of the imidazole compound of claim 1 or a pharmaceutically acceptable salt thereof for the production of an agent for the prophylaxis and/or treatment of impaired glucose tolerance disorder, diabetes, gestational diabetes, diabetic complications, insulin resistance syndrome, polycystic ovary syndrome, hyperlipidemia, atherosclerosis, cardiovascular diseases, hyperglycemia, pancreatitis, osteoporosis, hyperuricemia, hypertension, inflammatory bowel diseases, skin disorders related to an anomaly of differentiation of epidermic cells, angina pectoris, pulmonary hypertension, congestive heart failure, glomerulopathy, tubulointerstitial disorders, renal failure, angiostenosis, peripheral vascular diseases, cerebral apoplexy, chronic reversible obstructive impairment, autoimmune diseases, allergic rhinitis, urticaria, glaucoma, diseases characterized by impaired intestinal motility, impotence, nephritis, cancer cachexia, restenosis after PTCA, or cachexia.

10. A method of preventing and/or treating impaired glucose tolerance disorder, diabetes, gestational diabetes, diabetic complications, insulin resistance syndrome, polycystic ovary syndrome, hyperlipidemia, atherosclerosis, cardiovascular diseases, hyperglycemia, pancreatitis, osteoporosis, hyperuricemia, hypertension, inflammatory bowel diseases, skin disorders related to an anomaly of differentiation of epidermic cells, angina pectoris, pulmonary hypertension, congestive heart failure, glomerulopathy, tubulointerstitial disorders, renal failure, angiostenosis, peripheral vascular diseases, cerebral apoplexy, chronic reversible obstructive impairment, autoimmune diseases, allergic rhinitis, urticaria, glaucoma, diseases characterized by impaired intestinal motility, impotence,

(The following information was obtained from the records of the FBI, New York City Office, dated 7-10-68.)